- 11. The method of claim 1, wherein the molecular target comprises a tag moiety, and the aptamer is a high affinity binding partner to at least a portion of the tag moiety of the molecular target.
- 12. The method of claim 11, wherein the tag moiety is selected from a fluorescent protein or fragment thereof, a poly-His tag, a maltose binding protein tag, an albumin-binding protein tag, a calmodulin binding peptide tag, a glutathione S-transferase tag, a chitin binding protein tag, FLAG-tag, HA-tag, Protein A tag, and combinations thereof.
- 13. The method of claim 1, wherein the aptamer is selected from a non-naturally occurring aptamer and a naturally occurring aptamer.
- **14**. The method of claim **1**, wherein the aptamer is selected from a peptide aptamer, a DNA aptamer, and an RNA aptamer.
- **15**. The method of claim **1**, wherein said aptamer is an RNA aptamer comprising:
 - a core region comprising a nucleotide sequence of any one of SEQ ID NOs:1-13.
- **16**. The method of claim **1**, wherein the aptamer binds to a portion of a fluorescent protein selected from GFP, eGFP, CFP, eCFP, YFP, and eYFP.
- 17. The method of claim 1, wherein the aptamer is immobilized on a solid support.
- 18. The method of claim 1, wherein said molecular target comprises two or more associated biomolecules, and said analyzing comprises:
 - identifying each of the two or more associated biomolecules of the molecular target.
- 19. The method of claim 18, wherein said identifying comprises:
 - subjecting the molecular target to mass spectrometry to detect each of the two or more associated biomolecules of the molecular target.

- 20. The method of claim 1, wherein said analyzing comprises:
 - subjecting the molecular target to mass spectrometry to detect one or more posttranslational modifications of said molecular target.
- 21. The method of claim 20, wherein the one or more posttranslational modifications is selected from methylation, acetylation, phosphorylation, sumoylation, or combinations thereof.
- 22. The method of claim 1, wherein said molecular target comprises nucleotide oligomers and said analyzing comprises:
 - isolating the nucleotide oligomers from the molecular target and
 - subjecting said isolated nucleotide oligomers to an amplification reaction, a sequencing reaction, or a combination thereof to identify the isolated nucleotide oligomers of the molecular target.
- 23. The method of claim 1, wherein said analyzing comprises:
 - subjecting the molecular target to cryo-electron microscopy to determine the three-dimensional structure of the molecular target.
- 24. The method of claim 23, wherein said molecular target comprises two or more associated biomolecules, and said subjecting is carried out to determine the three-dimensional structure of the two or more associated biomolecules of the molecular target.
- 25. The method of claim 1, wherein the sample is a cell or tissue lysate, a plasma sample, a serum sample, a blood sample, an exosome sample, or other biological sample.

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